

replace paragraph on page 13, line 25 as follows:

Change(s) applied --(ii) recovering the multispecific antibody from the host  
to document, cell culture.--  
/K.D.D./

3/3/2011

Please replace the paragraph beginning at page 17, line 16 with the following rewritten paragraph:

--Figs. 2A-2C. Fig. 2A diagrams a selection scheme for C<sub>H</sub>3 heterodimer using phage display vector, pRA2. Phage displaying stable C<sub>H</sub>3 heterodimers are captured using an antibody directed to the gD flag. Fig. 2B diagrams a dicistronic operon in which C<sub>H</sub>3 expressed from a synthetic gene is co-secreted with a second copy of C<sub>H</sub>3 expressed from the natural gene (Ellison et al. Nucleic Acids Res. 10:4071-4079 (1982)) as a fusion protein with M13 gene III protein. The synthetic C<sub>H</sub>3 gene is preceded by a sequence encoding a peptide derived from herpes simplex virus glycoprotein D (gD flag, Lasky, L. A. and Dowbenko, D. J. (1984) DNA 3:23-29; Berman, P. W. et al., (1985) Science 227:1490-1492 and a cleavage (G) site for the site-specific protease, Genenase I (Carter, P. et al. (1989) Proteins: Structure, Function and Genetics 6:240-248). Fig. 2C is the nucleic acid sequence of the dicistronic operon (SEQ ID NO:13) of Fig. 2B in which the residues in the translated C<sub>H</sub>3 genes are numbered according to the Eu system of Kabat et al. In Sequences of Proteins of Immunological Interest, 5th ed. vol. 1, pp. 688-696, NIH, Bethesda, MD (1991). Protuberance mutation T366W is shown, as are the residues targeted for randomization in the natural C<sub>H</sub>3 gene (366, 368, and 407).--

Please replace the paragraph beginning at page 96, line 8 with the following rewritten paragraph:

--A large human single chain Fv (scFv) antibody library (Vaughan et al. (1996), *supra*) was panned for antibodies specific for eleven antigens including Axl(human receptor tyrosine kinase ECD), GCSF-R (human granulocyte colony stimulating factor receptor